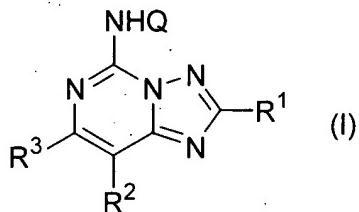


b.) Amendment to the Claims:

1. (Currently Amended) A [1,2,4]triazolo[1,5-c]pyrimidine derivative represented by formula (I):



{wherein

R<sup>1</sup> represents substituted or unsubstituted aryl, or a substituted or unsubstituted aromatic heterocyclic group;

R<sup>2</sup> represents a hydrogen atom, halogen, lower alkyl, lower alkanoyl, aroyl, substituted or unsubstituted aryl, or a substituted or unsubstituted aromatic heterocyclic group;

R<sup>3</sup> represents the following:

1) lower alkyl or hydroxy-substituted lower alkyl;

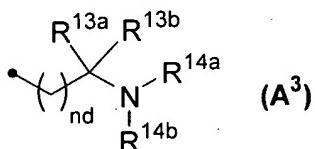
2) lower cycloalkyl;

3) formyl;

4) substituted or unsubstituted lower alkanoyl;

5) substituted or unsubstituted aroyl;

6) formula (A<sup>3</sup>)

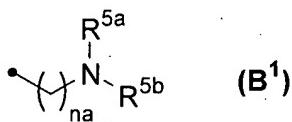


[wherein

nd represents an integer of 0 to 3;

$R^{13a}$  and  $R^{13b}$  may be the same or different and each represent a hydrogen atom, halogen, lower alkyl, lower cycloalkyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted aralkyl, substituted or unsubstituted aryl, a substituted or unsubstituted aromatic heterocyclic group, substituted or unsubstituted aroyl, lower alkoxy carbonyl, or lower alkoxy-substituted lower alkyl;  $R^{13a}$  and  $R^{13b}$  form a lower cycloalkane ring together with the adjacent carbon atom; or  $R^{13a}$  and  $R^{13b}$  are combined together to represent an oxygen atom or a sulfur atom; and

$R^{14a}$  and  $R^{14b}$  may be the same or different and each represent a hydrogen atom, substituted or unsubstituted lower alkyl, lower cycloalkyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted aryl, a substituted or unsubstituted aromatic heterocyclic group, substituted or unsubstituted aroyl, lower alkoxy carbonyl, formyl, or formula (B<sup>1</sup>)



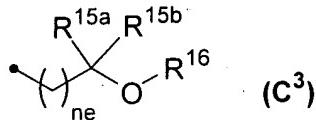
(wherein

$na$  represents an integer of 2 to 5; and

$R^{5a}$  and  $R^{5b}$  may be the same or different and each represent a hydrogen atom, lower alkyl, lower cycloalkyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted aralkyl, substituted or unsubstituted aryl, a substituted or unsubstituted aromatic heterocyclic group, substituted or unsubstituted aroyl, lower alkoxy carbonyl, lower alkoxy-substituted

lower alkyl, or formyl; or R<sup>5a</sup> and R<sup>5b</sup> form a substituted or unsubstituted heterocyclic group together with the adjacent nitrogen atom); or R<sup>14a</sup> and R<sup>14b</sup> form a substituted or unsubstituted heterocyclic group together with the adjacent nitrogen atom];

7) formula (C<sup>3</sup>)

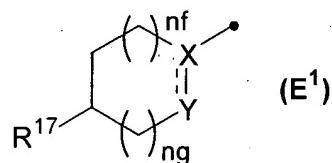


(wherein

ne, R<sup>15a</sup> and R<sup>15b</sup> have the same meanings as the above-described nd, R<sup>13a</sup> and R<sup>13b</sup>, respectively; and

R<sup>16</sup> represents a hydrogen atom, lower alkyl, lower cycloalkyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted aralkyl, substituted or unsubstituted aryl, a substituted or unsubstituted aromatic heterocyclic group, substituted or unsubstituted aroyl, or lower alkoxy-substituted lower alkyl);

8) formula (E<sup>1</sup>)



[wherein

nf represents an integer of 0 to 3;

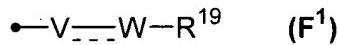
ng represents an integer of 1 to 4;

X---Y represents CR<sup>18</sup>-CH<sub>2</sub> (wherein R<sup>18</sup> represents a hydrogen atom, hydroxy,

halogen, nitro, cyano, trifluoromethyl, lower alkyl, lower alkoxy, lower alkanoyl, or lower alkoxycarbonyl), or C=CH; and

R<sup>17</sup> represents a hydrogen atom, substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted lower cycloalkyl, substituted or unsubstituted aryl, a substituted or unsubstituted aromatic heterocyclic group, substituted or unsubstituted aroyl, lower alkoxycarbonyl, or formyl];

9) formula (F<sup>1</sup>)

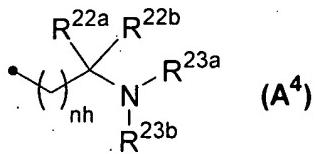


[wherein

V---W represents CR<sup>20</sup>=CR<sup>21</sup> (wherein R<sup>20</sup> and R<sup>21</sup> may be the same or different and each represent a hydrogen atom, substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted lower cycloalkyl, substituted or unsubstituted aryl, a substituted or unsubstituted aromatic heterocyclic group, substituted or unsubstituted aroyl, or lower alkoxycarbonyl) or C≡C; and

R<sup>19</sup> represents a hydrogen atom, substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted lower cycloalkyl, substituted or unsubstituted aryl, a substituted or unsubstituted aromatic heterocyclic group, substituted or unsubstituted aroyl, lower alkoxycarbonyl, or

formula (A<sup>4</sup>)



(wherein

*nh*,  $R^{22a}$ ,  $R^{22b}$ ,  $R^{23a}$  and  $R^{23b}$  have the same meanings as the above-

described *nd*,  $R^{13a}$ ,  $R^{13b}$ ,  $R^{14a}$  and  $R^{14b}$ , respectively),

provided that  $R^{19}$  is not substituted or unsubstituted aryl when  $V$ — $W$  is

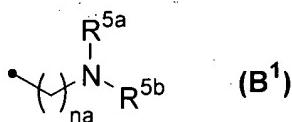
$\text{CH}=\text{CH}$ ];

10) aryl substituted with a substituent selected from the

group consisting of

$-\text{CH}_2\text{NHR}^{4a}$  [wherein  $R^{4a}$  represents substituted or unsubstituted lower alkyl, lower cycloalkyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted aryl, a substituted or unsubstituted aromatic heterocyclic group, substituted or unsubstituted aroyl, lower alkoxy carbonyl, formyl, or formula

(B<sup>1</sup>)

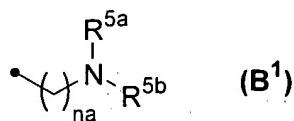


(wherein *na*,  $R^{5a}$  and  $R^{5b}$  have the same meanings as defined above,

respectively)],

$-(\text{CH}_2)_{nb}-\text{C}(\text{R}^{6a})(\text{R}^{6b})(\text{OR}^7)$  (wherein *nb*,  $R^{6a}$ ,  $R^{6b}$  and  $R^7$  have the same meanings as the above-described *nd*,  $R^{13a}$ ,  $R^{13b}$  and  $R^{16}$ , respectively), and  $-\text{NR}^{8a}\text{R}^{8b}$  [wherein  $\text{R}^{8b}$  and  $\text{R}^{8b}$  [wherein  $R^{8a}$  and  $R^{8b}$  may be the same or different and each represent a hydrogen atom, substituted or unsubstituted lower alkyl, lower cycloalkyl, substituted or unsubstituted lower alkanoyl,

substituted or unsubstituted aryl, a substituted or unsubstituted aromatic heterocyclic group, substituted or unsubstituted aroyl, lower alkoxycarbonyl, formyl, or formula (B<sup>1</sup>)



(wherein na, R<sup>5a</sup> and R<sup>5b</sup> have the same meanings as defined above, respectively)]; or

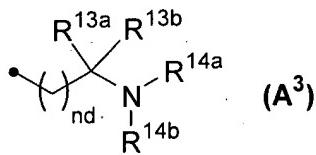
- 11) an aromatic heterocyclic group substituted with a substituent selected from the group consisting of -CH<sub>2</sub>NR<sup>4b</sup>R<sup>4c</sup> (wherein R<sup>4b</sup> and R<sup>4c</sup> have the same meanings as the above-described R<sup>14a</sup> and R<sup>14b</sup>, respectively), -(CH<sub>2</sub>)<sub>nb</sub>-C(R<sup>6a</sup>)(R<sup>6b</sup>)(OR<sup>7</sup>) (wherein nb, R<sup>6a</sup>, R<sup>6b</sup> and R<sup>7</sup> have the same meanings as defined above, respectively), and -NR<sup>8a</sup>R<sup>8b</sup> (~~wherein R<sup>8b</sup> and R<sup>8b</sup>~~ (wherein R<sup>8b</sup> and R<sup>8b</sup> have the same meanings as defined above, respectively); and

Q represents a hydrogen atom or 3,4-dimethoxybenzyl}, or a pharmaceutically acceptable salt thereof.

2. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 1, wherein R<sup>3</sup> is the following:

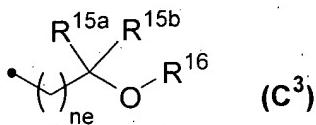
- 1) lower alkyl or hydroxy-substituted lower alkyl;
- 2) lower cycloalkyl;

- 3) formyl;
- 4) substituted or unsubstituted lower alkanoyl;
- 5) substituted or unsubstituted aroyl;
- 6) formula (A<sup>3</sup>)



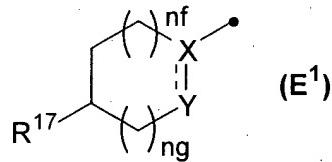
(wherein nd, R<sup>13a</sup>, R<sup>13b</sup>, R<sup>14a</sup> and R<sup>14b</sup> have the same meanings as defined above, respectively);

- 7) formula (C<sup>3</sup>)



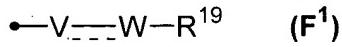
(wherein ne, R<sup>15a</sup>, R<sup>15b</sup> and R<sup>16</sup> have the same meanings as defined above, respectively);

- 8) formula (E<sup>1</sup>)



(wherein nf, ng, X---Y and R<sup>17</sup> have the same meanings as defined above, respectively); or

- 9) formula (F<sup>1</sup>)



(wherein V---W and R<sup>19</sup> have the same meanings as defined above, respectively),

or a pharmaceutically acceptable salt thereof.

3. (Currently Amended) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 1, wherein R<sup>3</sup> is aryl substituted with a substituent selected from the group consisting of

-CH<sub>2</sub>NHR<sup>4a</sup> (wherein R<sup>4a</sup> has the same meaning as defined above), -(CH<sub>2</sub>)<sub>nb</sub>-C(R<sup>6a</sup>)(R<sup>6b</sup>)(OR<sup>7</sup>) (wherein nb, R<sup>6a</sup>, R<sup>6b</sup> and R<sup>7</sup> have the same meanings as defined above, respectively), and -NR<sup>8a</sup>R<sup>8b</sup> (wherein R<sup>8b</sup> and R<sup>8b</sup> (wherein R<sup>8a</sup> and R<sup>8b</sup> have the same meanings as defined above, respectively), or a pharmaceutically acceptable salt thereof.

4. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 1, wherein R<sup>3</sup> is aryl substituted with -CH<sub>2</sub>NHR<sup>4a</sup> (wherein R<sup>4a</sup> has the same meaning as defined above), or a pharmaceutically acceptable salt thereof.

5. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 3 or 4, wherein the aryl is phenyl, or a pharmaceutically acceptable salt thereof.

6. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 1, wherein R<sup>3</sup> is an aromatic heterocyclic group substituted with a substituent selected from the group consisting of -CH<sub>2</sub>NR<sup>4b</sup>R<sup>4c</sup> (wherein R<sup>4b</sup> and R<sup>4c</sup> have the same meanings as defined above, respectively), -(CH<sub>2</sub>)<sub>nb</sub>-C(R<sup>6a</sup>)(R<sup>6b</sup>)(OR<sup>7</sup>) (wherein nb, R<sup>6a</sup>, R<sup>6b</sup> and R<sup>7</sup> have the same meanings as defined above, respectively), and -NR<sup>8a</sup>R<sup>8b</sup> (wherein R<sup>8b</sup> and R<sup>8b</sup> have the same meanings as defined above, respectively), or a pharmaceutically acceptable

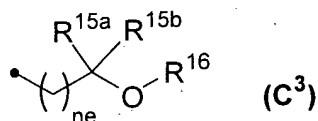
salt thereof.

7. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 1, wherein R<sup>3</sup> is an aromatic heterocyclic group substituted with -(CH<sub>2</sub>)<sub>nb</sub>-C(R<sup>6a</sup>)(R<sup>6b</sup>)(OR<sup>7</sup>) (wherein nb, R<sup>6a</sup>, R<sup>6b</sup> and R<sup>7</sup> have the same meanings as defined above, respectively), or a pharmaceutically acceptable salt thereof.

8. (Currently Amended) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 1, wherein R<sup>3</sup> is an aromatic heterocyclic group substituted with -NR<sup>8a</sup>R<sup>8b</sup> (wherein R<sup>8b</sup> and R<sup>8b</sup> (wherein R<sup>8a</sup> and R<sup>8b</sup> have the same meanings as defined above, respectively), or a pharmaceutically acceptable salt thereof.

9. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to any one of claims 6 to 8, wherein the aromatic heterocyclic group is pyridyl or thiazolyl, or a pharmaceutically acceptable salt thereof.

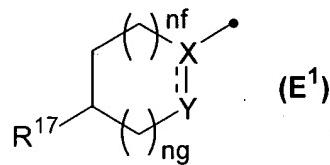
10. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 1, wherein R<sup>3</sup> is formula (C<sup>3</sup>)



(wherein ne, R<sup>15a</sup>, R<sup>15b</sup> and R<sup>16</sup> have the same meanings as defined above, respectively), or a pharmaceutically acceptable salt thereof.

11. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 1, wherein R<sup>3</sup> is -CH<sub>2</sub>OR<sup>16</sup> (wherein R<sup>16</sup> has the same meaning as defined above), or a pharmaceutically acceptable salt thereof.

12. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 1, wherein R<sup>3</sup> is formula (E<sup>1</sup>)

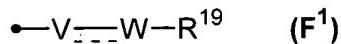


(wherein nf, ng, X---Y and R<sup>17</sup> have the same meanings as defined above, respectively), or a pharmaceutically acceptable salt thereof.

13. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 12, wherein nf is 1, ng is 1, and X---Y is C=CH, or a pharmaceutically acceptable salt thereof.

14. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 12 or 13, wherein R<sup>17</sup> is substituted or unsubstituted lower alkyl, or a pharmaceutically acceptable salt thereof.

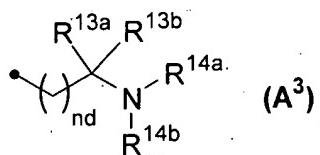
15. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 1, wherein R<sup>3</sup> is formula (F<sup>1</sup>)



(wherein V---W and R<sup>19</sup> have the same meanings as defined above, respectively), or a

pharmaceutically acceptable salt thereof.

16. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 1, wherein R<sup>3</sup> is formula (A<sup>3</sup>)



(wherein nd, R<sup>13a</sup>, R<sup>13b</sup>, R<sup>14a</sup> and R<sup>14b</sup> have the same meanings as defined above, respectively), or a pharmaceutically acceptable salt thereof.

17. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 16, wherein nd is 0, and R<sup>13a</sup> and R<sup>13b</sup> are combined together to represent an oxygen atom, or a pharmaceutically acceptable salt thereof.

18. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 16, wherein nd is 0, and R<sup>13a</sup> and R<sup>13b</sup> are each a hydrogen atom, or a pharmaceutically acceptable salt thereof.

19. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to any one of claims 16 to 18, wherein R<sup>14a</sup> and R<sup>14b</sup> may be the same or different and are each a hydrogen atom or substituted or unsubstituted lower alkyl, or a pharmaceutically acceptable salt thereof.

20. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to any

one of claims 16 to 18, wherein R<sup>14a</sup> and R<sup>14b</sup> form a substituted or unsubstituted heterocyclic group together with the adjacent nitrogen atom, or a pharmaceutically acceptable salt thereof.

21. (Original) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to claim 1, wherein R<sup>3</sup> is formyl, substituted or unsubstituted lower alkanoyl, or substituted or unsubstituted aroyl, or a pharmaceutically acceptable salt thereof.

22. (Currently Amended) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to any one of ~~claims 1 to 21~~, claims 1-4, 6-8, 10-13, 15-18 or 21, wherein Q is a hydrogen atom, or a pharmaceutically acceptable salt thereof.

23. (Currently Amended) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to ~~any one of claims 1 to 22~~, claim 22, wherein R<sup>1</sup> is furyl, or a pharmaceutically acceptable salt thereof.

24. (Currently Amended) The [1,2,4]triazolo[1,5-c]pyrimidine derivative according to ~~any one of claims 1 to 23~~, claim 23, wherein R<sup>2</sup> is a hydrogen atom, or a pharmaceutically acceptable salt thereof.

25. (Currently Amended) A pharmaceutical composition comprising the [1,2,4]triazolo[1,5-c]pyrimidine derivative according to ~~any one of claims 1 to 24~~ claim 22 or a pharmaceutically acceptable salt thereof as an active ingredient.

26. (Currently Amended) A therapeutic agent for Parkinson's disease comprising the [1,2,4]triazolo[1,5-c]pyrimidine derivative according to ~~any one of claims 1 to 24~~ claim 1 or a pharmaceutically acceptable salt thereof as an active ingredient.

27. (Currently Amended) A therapeutic agent for depression comprising the [1,2,4]triazolo[1,5-c]pyrimidine derivative according to ~~any one of claims 1 to 24~~ claim 1 or a pharmaceutically acceptable salt thereof as an active ingredient.

28. (Currently Amended) A therapeutic and/or preventive agent for a disease induced by hyperactivity of an adenosine A<sub>2A</sub> receptor, comprising the [1,2,4]triazolo[1,5-c]pyrimidine derivative according to ~~any one of claims 1 to 24~~ claim 1 or a pharmaceutically acceptable salt thereof as an active ingredient.

29. (Currently Amended) Use of the [1,2,4]triazolo[1,5-c]pyrimidine derivative according to ~~any one of claims 1 to 24~~ claim 1 or a pharmaceutically acceptable salt thereof for the manufacture of a therapeutic agent for Parkinson's disease.

30. (Currently Amended) Use of the [1,2,4]triazolo[1,5-c]pyrimidine derivative according to ~~any one of claims 1 to 24~~ claim 1 or a pharmaceutically acceptable salt thereof for the manufacture of a therapeutic agent for depression.

31. (Currently Amended) Use of the [1,2,4]triazolo[1,5-c]pyrimidine derivative

according to ~~any one of claims 1 to 24~~ claim 1 or a pharmaceutically acceptable salt thereof for the manufacture of a therapeutic and/or preventive agent for a disease induced by hyperactivity of an adenosine A<sub>2A</sub> receptor.

32. (Currently Amended) A therapeutic agent for a disease selected from the group consisting of Alzheimer's disease, progressive supranuclear palsy, AIDS encephalopathy, transmissible spongiform encephalopathy, multiple sclerosis, amyotrophic lateral sclerosis, Huntington's disease, multiple system atrophy, cerebral ischemia, sleep disorders, ischemic heart disease and intermittent claudications, comprising the [1,2,4]triazolo[1,5-c]pyrimidine derivative according to ~~any one of claims 1 to 24~~ claim 1 or a pharmaceutically acceptable salt thereof as an active ingredient.

33. (Currently Amended) Use of the [1,2,4]triazolo[1,5-c]pyrimidine derivative according to ~~any one of claims 1 to 24~~ claim 1 or a pharmaceutically acceptable salt thereof for the manufacture of a therapeutic agent for a disease selected from the group consisting of Alzheimer's disease, progressive supranuclear palsy, AIDS encephalopathy, transmissible spongiform encephalopathy, multiple sclerosis, amyotrophic lateral sclerosis, Huntington's disease, multiple system atrophy, cerebral ischemia, sleep disorders, ischemic heart disease and intermittent claudications.

34. (Currently Amended) A method for treating Parkinson's disease, comprising administering an effective amount of the [1,2,4]triazolo[1,5-c]pyrimidine derivative according to ~~any one of claims 1 to 24~~ claim 1 or a pharmaceutically acceptable salt thereof.

35. (Currently Amended) A method for treating depression, comprising administering an effective amount of the [1,2,4]triazolo[1,5-c]pyrimidine derivative according to ~~any one of claims 1 to 24~~ claim 1 or a pharmaceutically acceptable salt thereof.

36. (Currently Amended) A method for treating and/or preventing a disease induced by hyperactivity of an adenosine A<sub>2A</sub> receptor, comprising administering an effective amount of the [1,2,4]triazolo[1,5-c]pyrimidine derivative according to ~~any one of claims 1 to 24~~ claim 1 or a pharmaceutically acceptable salt thereof.

37. (Currently Amended) A method for treating a disease selected from the group consisting of Alzheimer's disease, progressive supranuclear palsy, AIDS encephalopathy, transmissible spongiform encephalopathy, multiple sclerosis, amyotrophic lateral sclerosis, Huntington's disease, multiple system atrophy, cerebral ischemia, sleep disorders, ischemic heart disease and intermittent claudications, comprising administering an effective amount of the [1,2,4]triazolo[1,5-c]pyrimidine derivative according to ~~any one of claims 1 to 24~~ claim 1 or a pharmaceutically acceptable salt thereof.